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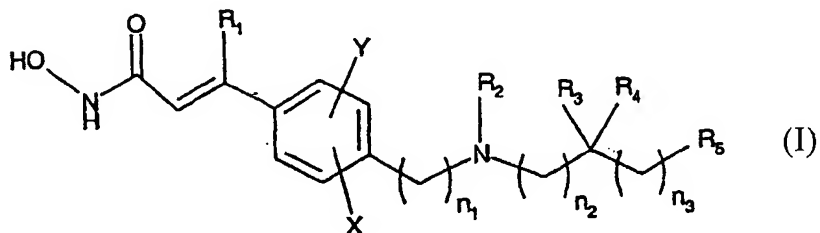
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(54) Title: HYDROXAMATE DERIVATIVES USEFUL AS DEACETYLASE INHIBITORS



(57) Abstract: The present invention provides hydroxamate compounds of formula (I) which are deacetylase inhibitors. The compounds are suitable for pharmaceutical compositions having anti-proliferative properties.

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INTERNATIONAL SEARCH REPORT

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A. CLASSIFICATION OF SUBJECT MATTER

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B. FIELDS SEARCHED

Minimum documentation searched (classification system followed by classification symbols)

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Documentation searched other than minimum documentation to the extent that such documents are included in the fields searched

Electronic data base consulted during the international search (name of data base and, where practical, search terms used)

EPO-Internal, CHEM ABS Data, BIOSIS, WPI Data, PAJ

C. DOCUMENTS CONSIDERED TO BE RELEVANT

Category *	Citation of document, with indication, where appropriate, of the relevant passages	Relevant to claim No.
X	WO 98 55449 A (QUEENSLAND INST MED RES ;FAIRLIE DAVID (AU); PARSONS PETER G (AU);) 10 December 1998 (1998-12-10) claims 4, 15 formula Vg, 16, 25-26; p. 6, l. 29-33; p. 13-p. 14, l. 11, p. 45 compound 5, p. 47 3rd structure, p. 48 MW2796, p. 65 table 1 ---	1-40
X	WO 95 31977 A (SLOAN KETTERING INST CANCER ;UNIV COLUMBIA (US)) 30 November 1995 (1995-11-30) claims 40, 42, 55-56; p. 23, l. 19-l. 35; p. 74 compound 77 and tables 2-3 --- -/--	1-40



Further documents are listed in the continuation of box C.



Patent family members are listed in annex.

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C.(Continuation) DOCUMENTS CONSIDERED TO BE RELEVANT

Category *	Citation of document, with indication, where appropriate, of the relevant passages	Relevant to claim No.
P,X	WO 01 38322 A (METHYLGENE INC) 31 May 2001 (2001-05-31) claims 26-27,29-31,37,41; compounds 36, 87-91,93-97,99-106, 108-115,120,145-147,170; table 5 ---	1-40
P,A	M. YOSHIDA ET AL.: "Histone deacetylase as a new target for cancer chemotherapy" CANCER CHEMOTHER. PHARMACOL., vol. 48, no. Suppl. 1, 1 July 2001 (2001-07-01), pages S20-S26, XP002188621 the whole document -----	1-40

INTERNATIONAL SEARCH REPORT

Information on patent family members

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